# PREFERRED DRUG LIST MEETING SCHEDULE

## State of Montana
### Department of Public Health & Human Services

## Montana Medicaid Drug Use Review Board/Formulary Committee Meeting

The State of Montana Medicaid Drug Utilization Review Board/Formulary Committee will tentatively hold a meeting on :

**Date:** April 5, 2006 (Wednesday)
**Time:** 1:00 pm – 4:00 pm Mountain Time
**Location:** AARP Montana Office 30 West 14th St, Suite 301, Helena, MT

At this time the Montana Medicaid Drug Utilization Review Board/Formulary Committee will tentatively review the following drug classes for Preferred Drug List ( PDL) review:

**Drug Class ReReviews**
**All drugs reviewed pertain to oral drugs unless otherwise indicated**

- ALPHA-GLUCOSIDASE INHIBITORS
- ANGIOTENSIN CONVERTING ENZYME INHIBITORS (ACEI) & COMBINATIONS
- ANGIOTENSIN II RECPTOR BLOCKERS (ARBS) & COMBINATIONS
- BETA-BLOCKERS-ORAL
- BISPHOSPHONATES
- INHALED CORTICOSTEROIDS
- INSULINS
- GLAUCOMA-PROSTAGLANDINS
- LIPOTROPICS-STATINS
- MEGLITINIDES
- NASAL CALCITONINS
- PROTON PUMP INHIBITORS
- THIAZOLIDIENEDIONES & COMBINATIONS
- TRIPTANS

**Public Testimony** will be taken into consideration in the committee's recommendations as to which drugs should be given preferred status in the above listed classes of medications for the state's Medicaid program. Sign-up for public comment will occur between 12:30pm -12:55 pm outside the Conference Room. See the General Procedures for Public Comment section of this document for further details

**Clinical Information**: Clinical information (in electronic format in PDF in the AMCP style dossier or desired style) may be sent on the drug classes listed above by March 22, 2006 to:

Mark Eichler, Mountain-Pacific Quality Health Foundation,
Tel: 406-457-5818, meichler@mpqhf.org and pdl@mt.gov

**Note:** This request constitutes a request for information pertaining to peer-reviewed literature including off label peer-reviewed studies or AMCP style - dossiers. Please note that all information sent is subject to public disclosure and that proprietary and confidential material should not be sent and that the sender accepts responsibility for all information sent. All information sent will be posted on a public website for viewing.

# Department of Public Health and Human Services
# DUR Board Meeting
# General Procedures for Public Comment

1. Thirty minutes prior to the beginning of the DUR Board Meeting, a sign up sheet for Public Comment will be posted for Pharmaceutical Manufacturers and Special Interest Groups for each Drug Class to be reviewed.

2. Sign up will close 5 minutes prior to the beginning of the DUR Board Meeting.

3. Speakers will be assigned on a first come basis respective to each Drug Class discussion.

4. Speakers will be asked to present **NEW INFORMATION ONLY** on their corresponding product or interest.

   a. New Information is considered the following: new product in the drug class, new indication since the last review or new studies released since the last review, excluding placebo only studies. New studies must be submitted in electronic format by March 22, 2006.

   b. Public comment will be allowed for up to 10 minutes to present new information about their product. However, please be respectful of your other colleagues and also of the Board's time. Please do not take 10 minutes if it is not needed. The DUR Board Coordinator has the option to end a speaker's comment time if the information is not relevant to the topic of discussion.

   c. Speakers must state their name, their affiliation, and whom they are speaking on behalf of or on request of, with any funding or payment agreements disclosed. Any studies cited during the presentation should be referenced with the primary source of funding included.

   d. Handouts are limited to two (2) pages (paper size: 8.5" by 11", one side only) of documentation. Access to computers or other technology presentation devices for slide presentations will not be available during this comment period.

   e. Public Comment will be limited to clinical and social comments; pricing or financial information regarding products and outcomes will not be permissible. The Board will be utilizing clinical information only. Information regarding pricing, cost or any other information of a financial nature will not be permissible and should not be discussed in handouts or during presentation by any public speaker.

   f. The speakers presenting handouts are asked to provide at least thirty (30) copies that will be distributed by Foundation staff to the DUR Board members, State staff and for public distribution.

   g. Copies will be collected by Foundation staff members at the time of sign-up.

   h. The State, FHSC and the DUR Board will be allowed to ask questions if needed during the presentation or after for clarification or discussion. Presenters will only be allowed to answer questions when specifically requested to do so by the Board during the remainder of the meeting.

   i. It is not permissible for presenters to interject or ask questions to DUR Board members during the session

5. Individual products may only be represented by one presentation. For example, a product jointly ventured by two pharmaceutical companies can only be represented once.

**Note: These procedures may be revised at the discretion of the Department.**